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CPC COOPERATIVE PATENT CLASSIFICATION

C CHEMISTRY; METALLURGY

(NOTES omitted)

CHEMISTRY

C01 INORGANIC CHEMISTRY

(NOTES omitted)

COMPOUNDS OF THE METALS BERYLLIUM, MAGNESIUM, ALUMINIUM, CALCIUM, STRONTIUM, BARIUM, RADIUM, THORIUM, OR OF THE RARE-

EARTH METALS (metal hydrides {monoborane, diborane or addition complexes thereof} C01B 6/00; salts of oxyacids of halogens C01B 11/00; peroxides, salts of peroxyacids C01B 15/00; sulfides or polysulfides of magnesium, calcium, strontium, or barium C01B 17/42; thiosulfates, dithionites, polythionates C01B 17/64; compounds containing selenium or tellurium C01B 19/00; binary compounds of nitrogen with metals C01B 21/06; azides C01B 21/08; {compounds other than ammonia or cyanogen containing nitrogen and non-metals and optionally metals C01B 21/082; amides or imides of silicon C01B 21/087}; metal {imides or} amides C01B 21/092, {C01B 21/0923}; nitrites C01B 21/50; {compounds of noble gases C01B 23/0005}; phosphides C01B 25/08; salts of oxyacids of phosphorus C01B 25/16; carbides C01B 32/90; compounds containing silicon C01B 33/00; compounds containing boron C01B 35/00; compounds having molecular sieve properties but not having base-exchange properties C01B 37/00; compounds having molecular sieve and base-exchange properties, e.g. crystalline zeolites, C01B 39/00; cyanides C01C 3/08; salts of cyanic acid C01C 3/14; salts of cyanamide C01C 3/16; thiocyanates C01C 3/20; {double sulfates of magnesium with sodium or potassium C01D 5/12; with other alkali metals C01D 15/00, C01D 17/00})

1/00	Methods of preparing compounds of the metals beryllium, magnesium, aluminium, calcium, strontium, barium, radium, thorium, or the rare earths, in general	 5/20 by precipitation from solutions of magnesium salts with ammonia 5/22 from magnesium compounds with alkali hydroxides or alkaline- earth oxides or
3/00 3/005 3/02	Compounds of beryllium • {Fluorides or double fluorides of beryllium with alkali metals or ammonium; Preparation of beryllium compounds therefrom} • Oxides; Hydroxides	hydroxides 5/24
5/00	Compounds of magnesium	chlorides, e.g. carnalite}
5/02 5/04 5/06 5/08 5/10 5/12	 Magnesia by oxidation of metallic magnesium by thermal decomposition of magnesium compounds (calcining magnesite or dolomite C04B 2/10) by calcining magnesium hydroxide by thermal decomposition of magnesium chloride with water vapour by thermal decomposition of magnesium sulfate, with or without reduction 	 5/32 . Preparation of anhydrous magnesium chloride by chlorinating magnesium compounds 5/34 . Dehydrating magnesium chloride containing water of crystallisation 5/36 . Bromides 5/38 . Magnesium nitrates 5/40 . Magnesium sulfates (double sulfates of magnesium with sodium or potassium C01D 5/12, with other alkali metals {C01D 15/00}, C01D 17/00) 5/42 . Magnesium sulfites
5/14 5/145 5/16	 Magnesium hydroxide {Purification} by treating magnesia, e.g. calcined dolomite, with water or solutions of salts not containing magnesium 	 7/00 Compounds of aluminium 7/02 Aluminium oxide; Aluminium hydroxide; Aluminates 7/021 • After-treatment of oxides or hydroxides 7/022 • Classification 7/023 • Grinding, deagglomeration or disintegration

7/025	Granulation or agglomeration	7/164	Calcium aluminates
7/025	Making or stabilising dispersions	7/164	Strontium aluminates
7/027	Treatment involving fusion or vaporisation	7/168	Barium aluminates
7/027	Beta-aluminas	7/18	Aluminium oxide or hydroxide from alkaline
7/04	Preparation of alkali metal aluminates;	7/10	earth metal aluminates
7704	Aluminium oxide or hydroxide therefrom	7/20	Preparation of aluminium oxide or hydroxide
	(C01F 7/028 takes precedence)		from aluminous ores using acids or salts
7/043	Lithium aluminates	7/22	with halides or halogen acids
7/046	Stabilisation of aluminates	7/24	with nitric acid or nitrogen oxides
7/06	by treating aluminous minerals or waste-	7/26	with sulfuric acids or sulfates
	like raw materials with alkali hydroxide,	7/28	with sulfurous acid
	e.g. leaching of bauxite according to the	7/30	Preparation of aluminium oxide or hydroxide
	Bayer process (obtaining aluminium oxide or		by thermal decomposition or by hydrolysis or
	hydroxide from the resulting aluminate solution		oxidation of aluminium compounds
7/0/0/	C01F 7/14)	7/302	Hydrolysis or oxidation of gaseous aluminium
7/0606	Making-up the alkali hydroxide solution		compounds in the gaseous phase
7/0613	from recycled spent liquor Pretreatment of the minerals, e.g. grinding	7/304	of organic aluminium compounds
7/0613	Digestion	7/306	Thermal decomposition of hydrated chlorides,
7/0626	Processes making use of tube digestion	7/200	e.g. of aluminium trichloride hexahydrate
7/0020	only	7/308	Thermal decomposition of nitrates
7/0633	characterised by the use of additives	7/32	Thermal decomposition of sulfates including
7/064	Apparatus for digestion, e.g. digestor	7/34	complex sulfates, e.g. alums • Preparation of aluminium hydroxide by
77004	vessels or heat exchangers	1/34	precipitation from solutions containing
7/0646	• • • • Separation of the insoluble residue, e.g. of		aluminium salts
	red mud	7/36	from organic aluminium salts
7/0653	characterised by the flocculant added	7/38	Preparation of aluminium oxide by thermal
	to the slurry (final clarification of the	1150	reduction of aluminous minerals
	aluminate solution C01F 7/47)	7/40	in the presence of aluminium sulfide
7/066	Treatment of the separated residue	7/42	Preparation of aluminium oxide or hydroxide
7/0666	Process control or regulation		from metallic aluminium, e.g. by oxidation
7/0673	from phosphate-containing minerals	7/422	by oxidation with a gaseous oxidator at a high
7/068	• • • from carbonate-containing minerals, e.g.		temperature
	dawsonite	7/424	using a plasma
7/0686	from sulfate-containing minerals, e.g. alunite	7/426	by applying mechanical energy to solid
7/0693	from waste-like raw materials, e.g. fly ash or		aluminium at a low temperature
7.00	Bayer calcination dust	7/428	• • • by oxidation in an aqueous solution
7/08	• • by treating aluminous minerals with sodium	7/44	• Dehydration of aluminium oxide or hydroxide,
	carbonate, e.g. sinter processes (C01F 7/0613, C01F 7/066 take precedence)		i.e. all conversions of one form into another
7/085	according to the lime-sinter process	7/441	involving a loss of water
7/10	by treating aluminous minerals with alkali	7/441	by calcination
7/10	sulfates and reducing agents	7/442	in presence of a calcination additive
7/12	Alkali metal aluminates from alkaline-earth	7/444	Apparatus therefor
7712	metal aluminates	7/445	making use of a fluidised bed
7/14	Aluminium oxide or hydroxide from alkali	7/447	by wet processes
	metal aluminates	7/448	 using superatmospheric pressure, e.g. hydrothermal conversion of gibbsite into
7/141	from aqueous aluminate solutions by		boehmite
	neutralisation with an acidic agent	7/46	Purification of aluminium oxide, aluminium
7/142	• • • • with carbon dioxide	77 10	hydroxide or aluminates (C01F 7/028 takes
7/144	• • • from aqueous aluminate solutions by		precedence)
	precipitation due to cooling, e.g. as part of	7/47	of aluminates, e.g. removal of compounds of
	the Bayer process		Si, Fe, Ga or of organic compounds from Bayer
7/145	characterised by the use of a crystal		process liquors
	growth modifying agent other than	7/473	Removal of organic compounds, e.g. sodium
7/1/7	aluminium hydroxide seed		oxalate
7/147 7/148	 Apparatus for precipitation Separation of the obtained hydroxide, e.g.	7/476	by oxidation
//148	by filtration or dewatering	7/48	Halides, with or without other cations besides
7/16	Preparation of alkaline-earth metal aluminates	= 15°	aluminium
,,10	or magnesium aluminates; Aluminium oxide	7/50	. Fluorides
	or hydroxide therefrom (C01F 7/028 takes	7/52	Double compounds containing both fluorine
	precedence)		and other halide groups
7/162	Magnesium aluminates		

7/54	• • • Double compounds containing both aluminium and alkali metals or alkaline-earth metals	11/181	• • {Preparation of calcium carbonate by carbonation of aqueous solutions and characterised by control
7/56	• • Chlorides (containing fluorine <u>C01F 7/52</u>)		of the carbonation conditions}
7/57	• • Basic aluminium chlorides, e.g. polyaluminium chlorides	11/182	• • {Preparation of calcium carbonate by carbonation of aqueous solutions and characterised by an
7/58	Preparation of anhydrous aluminium chloride		additive other than CaCO ₃ -seeds}
7/60	from oxygen-containing aluminium	11/183	{the additive being an organic compound}
	compounds	11/184	• • {Preparation of calcium carbonate by carbonation
7/62	Purification		of solutions based on non-aqueous solvents}
7/64	• • Bromides (containing fluorine <u>C01F 7/52</u>)	11/185	• • {After-treatment, e.g. grinding, purification,
7/66	• Nitrates, with or without other cations besides		conversion of crystal morphology}
7700	aluminium	11/186	• • {Strontium or barium carbonate}
7/68	Aluminium compounds containing sulfur	11/187	{Strontium carbonate}
7/70	Sulfides	11/188	{Barium carbonate}
7/72	. Sulfites	11/20	• Halides
7/74	. Sulfates	11/22	Fluorides
7/741	Preparation from elemental aluminium or	11/24	• Chlorides
///41	elemental aluminium containing materials, e.g.	11/26	from sulfides
	foil or dross	11/28	by chlorination of alkaline-earth metal
7/743	Preparation from silicoaluminious materials,	11/20	compounds
1/173	e.g. clays or bauxite	11/30	Concentrating; Dehydrating; Preventing the
7/745	• • • Preparation from alums, e.g. alunite	11/30	adsorption of moisture or caking
7/746	After-treatment, e.g. dehydration or	11/32	Purification
77740	stabilisation	11/34	. Bromides
7/748	Purification	11/34	Nitrates
7/76	Double salts, i.e. compounds containing,	11/38	Preparation with nitric acid or nitrogen oxides
7/70	besides aluminium and sulfate ions, only other	11/38	
	cations, e.g. alums	11/40	Preparation by double decomposition with nitrates
7/762	Ammonium or alkali metal aluminium	11/42	 Double salts (with magnesium <u>C01F 5/38</u>)
11102	sulfates	11/42	Concentrating; Crystallisating; Dehydrating;
7/765	Ammonium aluminium sulfates	11/44	Preventing the absorption of moisture or caking
7/767	Alkaline earth metal aluminium sulfates	11/46	Sulfates (dehydration of gypsum {for the production
7/77	Aluminium carbonates	11/40	of calcium sulfate cements \(\) \(\frac{C04B 11/02}{C04B 11/02} \)
7/78	Compounds containing aluminium and two or	11/462	• • {Sulfates of Sr or Ba}
1/10	more other elements, with the exception of oxygen	11/462	• {Sulfates of St of Ba}• {Sulfates of Ca from gases containing sulfur
	and hydrogen (aluminates C01F 7/02; compounds	11/404	oxides}
	containing aluminium, fluorine and alkali or	11/466	• • {Conversion of one form of calcium sulfate to
	alkaline earth metals <u>C01F 7/54</u> ; nitrates containing	11/400	another}
	other cations besides aluminium C01F 7/66;	11/468	• • {Purification of calcium sulfates}
	sulfides, sulfites or sulfates containing other cations	11/48	Sulfites
	besides aluminium <u>C01F 7/70</u> - <u>C01F 7/74</u>)	11/40	• Bunics
7/782	containing carbonate ions, e.g. dawsonite	13/00	Compounds of radium
7/784	. Layered double hydroxide, e.g. comprising	15/00	Compounds of thorium
	nitrate, sulfate or carbonate ions as intercalating	15/00	Compounds of thortum
	anions	17/00	Compounds of rare earth metals
7/785	Hydrotalcite		NOTES
7/786	containing, besides aluminium, only anions, e.g.		
	$Al[OH]_xCl_y[SO_4]_z$ (mixed halides $C01F7/48$)		1. In this group, the following expression is used
7/788	Ammonium aluminium fluorides, e.g. ammonium		with the meaning indicated; "rare earth metals"
	hexafluoroaluminate		means elements from the group of the lanthanides
11/00	Compounds of calcium, strontium, or barium		as well as scandium or yttrium, taken alone or in combination.
11/00	(C01F 7/00 takes precedence)		2. When classifying a compound in groups
11/005	• {Preparation involving liquid-liquid extraction,		Colf 17/20 - Colf 17/38, then its specific
11/005	absorption or ion-exchange}		preparation or treatment must also be classified in
11/02	Oxides or hydroxides (production of lime		groups $C01F 17/10 - C01F 17/17$ as long as the
/ 02	C04B 2/00)		compound is characterised by its preparation or
11/04	by thermal decomposition		treatment and vice versa.
11/06	of carbonates		
11/08	by reduction of sulfates	17/10	• Preparation or treatment, e.g. separation or
11/00	from sulfides		purification
11/10	from silicates	17/13	• • by using ion exchange resins, e.g. chelate resins
11/12		17/17	 involving a liquid-liquid extraction
	Purification		
11/18	. Purification. Carbonates		

17/20	Compounds containing only rare earth metals as the metal element
17/206	oxide or hydroxide being the only anion
17/212	Scandium oxides or hydroxides
17/212	•
	Yttrium oxides or hydroxides
17/224	Oxides or hydroxides of lanthanides
17/229	Lanthanum oxides or hydroxides
17/235	Cerium oxides or hydroxides
17/241	• • • containing two or more rare earth metals, e.g. NdPrO ₃ or LaNdPrO ₃
17/247	Carbonates
17/253	Halides
17/259	Oxyhalides
17/265	Fluorides
17/271	Chlorides
17/276	Nitrates
17/282	Sulfates
17/288	Sulfides
17/294	Oxysulfides
17/30	. Compounds containing rare earth metals and
	at least one element other than a rare earth
	metal, oxygen or hydrogen, e.g. La ₄ S ₃ Br ₆
	(<u>C01F 17/247</u> - <u>C01F 17/294</u> take precedence)
17/32	• oxide or hydroxide being the only anion, e.g.
	NaCeO ₂ or Mg _x Ca _y EuO
17/34	Aluminates, e.g. YAlO ₃ or Y _{3-x} Gd _x Al ₅ O ₁₂
17/36	halogen being the only anion, e.g. NaYF ₄
17/38	sulfur being the only anion, e.g. CaLa ₂ S ₄